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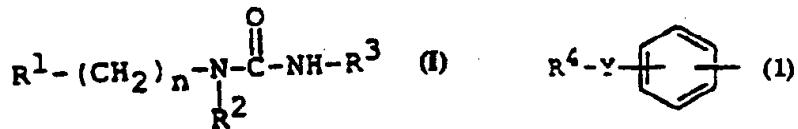
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(54) Title: UREA DERIVATIVES AND THEIR USE AS ACAT-INHIBITORS

(57) Abstract

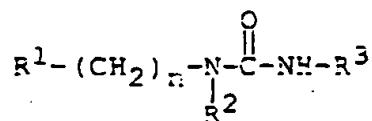
Urea derivatives of formula (I), wherein R¹ is a group of formula (I) (in which R⁴ is aryl which may have suitable substituent(s), or heterocyclic group which may have suitable substituent(s), and Y is bond, lower alkylene, -S-, -O-, (a), -CH-, -CONH-, (b), (in which R⁷ is lower alkyl), -NHSO₂-, -SO₂NH-, -SO₂NHCO- or -CONHSO₂-); or thiazolyl, imidazolyl, pyrazolyl, pyridyl, thiienyl, furyl, isoxazolyl or chromanyl, each of which may have suitable substituent(s); R² is lower alkyl, lower alkoxy(lower)alkyl, cycloalkyl, ar(lower)alkyl which may have suitable substituent(s), heterocyclic group or heterocyclic(lower)alkyl, R³ is aryl which may have suitable substituent(s) or heterocyclic group which may have suitable substituent(s), and n is 0 or 1, and a pharmaceutically acceptable salt thereof which are useful as a medicament in the treatment of hypercholesterolemia, hyperlipidemia and atherosclerosis.



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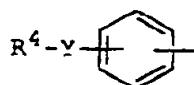
C L A I M S

1. A compound of the formula :



wherein

R^1 is a group of the formula :



(in which

R^4 is aryl which may have suitable substituent(s), or heterocyclic group which may have suitable substituent(s); and

C

||

y is bond, lower alkylene, -S-, -O-, -C-,
 $=CH-$, -CONH-, -N-CO-, (in which R^7 is lower
 R^7 alkyl),
 $-NHSO_2-$, $-SO_2NH-$, $-SO_2NHCO-$ or $-CONHSO_2-$);
 or

thiazolyl, imidazolyl, pyrazolyl, pyridyl,
 thienyl, furyl, isoxazolyl or chromanyl, each of
 which may have suitable substituent(s);

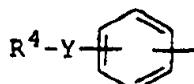
R^2 is lower alkyl, lower alkoxy(lower)alkyl,
 cycloalkyl, ar(lower)alkyl which may have
 suitable substituent(s), heterocyclic group or
 heterocyclic(lower)alkyl,

R^3 is aryl which may have suitable substituent(s) or
 heterocyclic group which may have suitable

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substituent(s), and
n is 0 or 1,
and a pharmaceutically acceptable salt thereof.

5 2. A compound of claim 1, wherein
R¹ is a group of the formula :



(in which
15 R⁴ is phenyl which may have 1 to 3 substituent(s)
selected from the group consisting of
halogen, lower alkyl, di(lower)alkylamino,
protected amino, cyano, heterocyclic group
which may have mono(or di or tri)-
ar(lower)alkyl, hydroxy, protected hydroxy
and mono(or di or tri)halo(lower)alkyl;
20 or thienyl, pyrazolyl, imidazolyl,
triazolyl, pyridyl, pyrrolyl, tetrazolyl,
oxazolyl, thiazolyl, oxadiazolyl,
piperazinyl, thiazolidinyl or
methylenedioxyphenyl, each of which may have
25 1 to 3 substituent(s) selected from the
group consisting of lower alkyl, mono(or di
or tri)ar(lower)alkyl and oxo;

30 Y is bond, lower alkylene, -S-, -O-, -C-, =CH-,
-CONH-, -N-CO- (in which R⁷ is lower alkyl),
R⁷
-NHSO₂-, -SO₂NH-, -SO₂NHCO- or -CONHSO₂-);
or
35 thiazolyl, imidazolyl, pyrazolyl, pyridyl,